Section: Contact Information
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Section: Project Requirements and Description

Group: Requirements to submit AOI
A comprehensive review of previously published data has been completed. : Yes
The specific aims are clear and focused. : Yes
The investigator has appropriate experience and expertise to develop the concept proposal; if not, has identified a mentor or senior co-investigator. : Yes
The investigator agrees to develop an initial draft of the concept proposal within 6 weeks of approval of the AOI and to finalize the concept proposal within 6 months. : Yes
Project Title: Neurocognitive functioning, emotional and quality of life of childhood cancer survivors of Asian/Pacific Islanders
Planned research population (eligibility criteria):
I would like to see roughly 400 Asian/Pacific Islanders in the CCSS Cohort as a descriptive study.
Proposed specific aims:
Aim 1: To characterize neurocognitive, emotional, quality of life and social attainment outcomes in cancer survivors of Asian/Pacific islander descent in the CCSS cohort.
Aim 2: To compare neurocognitive, emotional, quality of life and social attainment outcomes between cancer survivors of Asian/Pacific islander descent and caucasian descent, adjusting for diagnosis and demographic characteristics.
Aim 3: To identify demographic and treatment predictors of neurocognitive, emotional and quality of life outcomes in cancer survivors of Asian/Pacific islander descent.

Will the project require non-CCSS funding to complete? : No
If yes, what would be the anticipated source(s) and timeline(s) for securing funding?:

Group: Does this project require contact of CCSS study subjects for:
Additional self-reported information: No
Biological samples: No
Medical record data: No
If yes to any of the above, please briefly describe.

**Group: What CCSS Working Group(s) would likely be involved? (Check all that apply)**
- Second Malignancy:
- Chronic Disease:
- Psychology / Neuropsychology: Primary
- Genetics:
- Cancer Control:
- Epidemiology / Biostatistics:

**Section: Outcomes or Correlative Factors**
- Late mortality:
- Second Malignancy:

**Group: Health Behaviors**
- Tobacco:
- Alcohol:
- Physical activity:
- Medical screening:
- Other:
  If other, please specify:

**Group: Psychosocial**
- Insurance: Primary
- Marriage: Primary
- Education: Primary
- Employment: Primary
- Other:
  If other, please specify:

**Group: Medical Conditions**
- Hearing/Vision/Speech:
- Hormonal systems:
- Heart and vascular:
- Respiratory:
- Digestive:
- Surgical procedures:
- Brain and nervous system:
- Other:
If other, please specify:

**Group: Medications**
Describe medications:

**Group: Psychologic/Quality of Life**
BSI-18: Primary
SF-36: Primary
CCSS-NCQ: Primary
PTS :
PTG :
Other :
If other, please specify :

**Group: Other**
Pregnancy and offspring :
Family history :
Chronic conditions (CTCAE v3) :
Health status :

**Group: Demographic**
Age: Primary
Race: Primary
Sex: Primary
Other :
If other, please specify :

**Group: Cancer treatment**
Chemotherapy: Correlative Factors
Radiation therapy: Correlative Factors
Surgery :

**Section: Anticipated Sources of Statistical Support**
CCSS Statistical Center: Yes
Local institutional statistician :
If local, please provide the name(s) and contact information of the statistician(s) to be involved. :
Will this project utilize CCSS biologic samples? : No
If yes, which of the following? :
If other, please explain :

**Section: Other General Comments**
Other General Comments :